

University of Dundee

Hypoxia is not the primary mechanism contributing to exercise-induced proteinuria

Joyce, Kelsley E.; Delamere, John; Bradwell, Susie; Myers, Stephen David; Ashdown, Kimberly; Rue, Carla

Published in:
BMJ Open Sport and Exercise Medicine

DOI:
[10.1136/bmjsem-2019-000662](https://doi.org/10.1136/bmjsem-2019-000662)

Publication date:
2020

Licence:
CC BY-NC

Document Version
Publisher's PDF, also known as Version of record

[Link to publication in Discovery Research Portal](#)

Citation for published version (APA):

Joyce, K. E., Delamere, J., Bradwell, S., Myers, S. D., Ashdown, K., Rue, C., Lucas, S. J. E., Thomas, O. D., Fountain, A., Edsell, M., Myers, F., Malein, W., Imray, C., Clarke, A., Lewis, C. T., Newman, C., Johnson, B., Cadigan, P., Wright, A., & Bradwell, A. (2020). Hypoxia is not the primary mechanism contributing to exercise-induced proteinuria. *BMJ Open Sport and Exercise Medicine*, 6(1), 1-10. [e000662].
<https://doi.org/10.1136/bmjsem-2019-000662>

General rights



Copyright and moral rights for the publications made accessible in Discovery Research Portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

- Users may download and print one copy of any publication from Discovery Research Portal for the purpose of private study or research.
- You may not further distribute the material or use it for any profit-making activity or commercial gain.
- You may freely distribute the URL identifying the publication in the public portal.

Take down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Hypoxia is not the primary mechanism contributing to exercise-induced proteinuria

Kelsley E Joyce ^{1,2}, John Delamere,^{2,3} Susie Bradwell,^{2,4} Stephen David Myers,^{2,5} Kimberly Ashdown,^{2,5} Carla Rue,^{2,5} Samuel JE Lucas,^{1,2} Owen D Thomas,^{2,3} Amy Fountain,⁶ Mark Edsell,^{2,7} Fiona Myers,⁸ Will Malein,^{2,9} Chris Imray,^{2,10} Alex Clarke,^{2,11} Christopher T Lewis,^{2,12} Charles Newman,^{2,13} Brian Johnson,^{2,14} Patrick Cadigan,² Alexander Wright,^{2,15} Arthur Bradwell ^{2,3}

To cite: Joyce KE, Delamere J, Bradwell S, *et al.* Hypoxia is not the primary mechanism contributing to exercise-induced proteinuria. *BMJ Open Sport & Exercise Medicine* 2020;**6**:e000662. doi:10.1136/bmjsem-2019-000662

► Additional material is published online only. To view please visit the journal online (<http://dx.doi.org/10.1136/bmjsem-2019-000662>).

Accepted 23 February 2020

ABSTRACT

Introduction Proteinuria increases at altitude and with exercise, potentially as a result of hypoxia. Using urinary alpha-1 acid glycoprotein (α 1-AGP) levels as a sensitive marker of proteinuria, we examined the impact of relative hypoxia due to high altitude and blood pressure-lowering medication on post-exercise proteinuria.

Methods Twenty individuals were pair-matched for sex, age and ACE genotype. They completed maximal exercise tests once at sea level and twice at altitude (5035 m). Losartan (100 mg/day; angiotensin-receptor blocker) and placebo were randomly assigned within each pair 21 days before ascent. The first altitude exercise test was completed within 24–48 hours of arrival (each pair within ~1 hour). Acetazolamide (125 mg two times per day) was administered immediately after this test for 48 hours until the second altitude exercise test.

Results With placebo, post-exercise α 1-AGP levels were similar at sea level and altitude. Odds ratio (OR) for increased resting α 1-AGP at altitude versus sea level was greater without losartan (2.16 times greater). At altitude, OR for reduced post-exercise α 1-AGP (58% lower) was higher with losartan than placebo (2.25 times greater, $p=0.059$) despite similar pulse oximetry (SpO_2) ($p=0.95$) between groups. Acetazolamide reduced post-exercise proteinuria by approximately threefold (9.3 ± 9.7 vs 3.6 ± 6.0 μ g/min; $p=0.025$) although changes were not correlated ($r=-0.10$) with significant improvements in SpO_2 ($69.1\pm 4.5\%$ vs $75.8\pm 3.8\%$; $p=0.001$).

Discussion Profound systemic hypoxia imposed by altitude does not result in greater post-exercise proteinuria than sea level. Losartan and acetazolamide may attenuate post-exercise proteinuria, however further research is warranted.

INTRODUCTION

Proteinuria typically results from protein leakage from the capillary lumen through the glomerular filter,¹ with some removal in the tubules, as shown by studies inhibiting renal tubular reabsorption with lysine infusions.² Comparisons between albumin, a selectively reabsorbed,³ 66 kDa, negatively charged (pI

Summary box

- Post-exercise proteinuria is not directly related to systemic hypoxia but it is related to exercise intensity.
- Losartan may attenuate post-exercise proteinuria by maintaining the charge-selectivity function in the glomerular filter, although further investigations are warranted.
- Future proteinuria research should aim to include urinary alpha-1 acid glycoprotein and it should consider analyses using automated turbidimetric immunoassays.

4.7) protein that passes through the glomerular membrane via the slit diaphragm pores,⁴ and alpha-1 acid glycoprotein (α 1-AGP),^{5–7} a smaller (41–43 kDa) and more negatively charged protein (pI 2.7)^{5–7} indicate that urinary α 1-AGP is a more sensitive marker of glomerular leakage than albumin.^{7–9} Urinary α 1-AGP excretion has implicated the glomerular origin of the proteinuria exhibited with both, altitude⁷ and post-exercise.⁵ Hypothesised mechanisms for such glomerular leak have included changes in renal blood flow,^{5–10} increases in peritubular pressure and blood pressure² (BP), hypoxia¹¹ and acid–base disturbances,² although the mechanisms remain unclear.¹² The contributions of BP and hypoxia, especially to post-exercise proteinuria may be uniquely evaluated with altitude exercise.

At altitude, exercise BP is amplified^{13–14} and such exaggerations are responsive to anti-hypertensive therapies such as angiotensin II type 1 (AT1) receptor antagonists or blockers (ARBs).^{13–15} ARBs reduce BP via several mechanisms, including selectively blocking angiotensin II (A-II) from binding to AT1s within the vasculature,¹⁶ promoting vasodilation. Specific to the kidney, ARBs



© Author(s) (or their employer(s)) 2020. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Kelsley E Joyce;
kej764@student.bham.ac.uk

(eg, losartan) 'block' vasoconstriction imposed by A-II on afferent arterioles.¹⁷ ARBs also limit both, vasopressin secretion and aldosterone production, further adding to its BP lowering effect.¹⁶ Considering these factors, it would be reasonable to expect altitude post-exercise proteinuria to be greater than at sea level and that AT1 blockers such as losartan could attenuate the post-exercise response. To evaluate this, we planned to compare post-exercise α 1-AGP excretion between (1) sea level and altitude exercise and (2) losartan and placebo groups following exercise at altitude.

Exercise oxygen saturation is also altered at altitude and it is profoundly lower compared with sea level¹⁸ but it can be improved by acetazolamide,¹⁹ a carbonic anhydrase inhibitor commonly used to prevent altitude illness.^{20 21} By inhibiting carbonic anhydrase, a catalyst of the reversible CO₂ hydration reaction, acetazolamide promotes carbonic acid formation and dissociation of H⁺ and bicarbonate in the blood²¹ which limits hypoxia-induced alkalosis and improves oxygen saturation.²¹ Assuming hypoxia as the main mechanism of post-exercise proteinuria, altitude exercise would be expected to produce larger increases in post-exercise proteinuria compared with sea level. In addition, subsequent improvements in arterial oxygenation would be expected to have the 'reverse' effect. To evaluate this, we planned to compare post-exercise α 1-AGP between (1) sea level and altitude and (2) placebo and placebo +acetazolamide groups at altitude.

We hypothesised that exercise at altitude would increase post-exercise α 1-AGP levels compared with sea level, primarily due to greater systemic hypoxia, and that losartan and acetazolamide would attenuate the observed increases by improving blood and peritubular pressures¹⁶ and alleviating renal hypoxia,²² respectively.

As part of an expedition to Quito, Ecuador, we measured post-exercise urinary α 1-AGP excretion in 20 pair-matched individuals once at sea level (before placebo or losartan administration) and twice at altitude as outlined in the following sections.

METHODS

Design and participants

Twenty participants (14 men, 6 women) free of any pre-existing conditions were included in the study. ACE genotyping (II, ID or DD) was performed to limit potential differences between groups that could be attributed to ACE genotype (eg, response to losartan,²³ response to altitude,²⁴ etc.). Participants were pair-matched participants for ACE genotype, age, sex, previous altitude exposure and glomerular filtration rate (GFR, obtained 4 weeks prior to ascent) (figure 1).

Following matching, a double-blind, randomised, placebo-controlled trial design was adopted. Individuals within each pair were randomly assigned to either placebo or losartan group (figure 1). Losartan administration (100mg/day or placebo) began in the UK 21 days prior to departure for Quito, Ecuador (2850m). On

arrival at Quito, participants ascended over 10 days to the Whymper Hut on the flank of Chimborazo volcano (5035m, figure 1).

Baseline and daily measures

Baseline measures of height, body mass, GFR and creatinine (μ mol/L) were recorded 4 weeks prior to ascent (figure 1A). Estimated GFR (eGFR) was calculated for each individual using the Modification of Diet in Renal Disease study equation²⁵:

$$\text{eGFR}(\text{mL}/\text{min}/1.73\text{m}^2) = 32\,788 \times (\text{serum creatinine } \mu\text{mol}/\text{litre})^{-1.154} \times (\text{Age})^{-0.203} \text{ (then } 0.742 \text{ for females)}$$

During ascent (figure 1B), resting measures of systolic BP (SBP) and diastolic BP (DBP) were collected each morning using a manual sphygmomanometer.

Exercise protocols and measures

Sea-level exercise

Baseline sea-level graded maximal exercise tests were performed 4 weeks prior to ascent on a cycle ergometer (Alticycle) designed for altitude studies²⁶ (figure 1A). Volitional fatigue was used as the primary end point criterion for the test.²⁷ Maximal power output (Watt_{max}) was measured by the Alticycle and heart rate (HR) recordings were facilitated by telemetry (Polar Electro, UK).

Altitude exercise

The first altitude exercise tests were commenced on arrival to the Whymper Hut for five pairs (day 7) and completed on the following day (day 8) for remaining pairs (figure 1). These initial tests were immediately followed by acetazolamide administration (125mg orally, two times per day) which occurred 48 hours prior to the second round of altitude exercise tests in all participants. Similar to the first tests, the second round of altitude exercise tests were performed across 2 days (on days 9 and 10, figure 1).

At altitude, pre-exercise measurements of oxygen uptake (VO₂), carbon dioxide production (VCO₂), ventilation (V_E), HR, SBP and DBP were collected. Participants undertook a 5 min self-paced warm-up followed by a modified graded exercise test on the Alticycle which was commenced at 30% of sea-level Watt_{max}. Intensity was increased every 3 min by 10% until the participant reached 80% of sea-level Watt_{max}. From this point, intensity was increased by 10% each minute until volitional fatigue. Expired respiratory gases were analysed breath-by-breath using a Cosmed K4b² (Metabolic Company, Rome, Italy) portable metabolic system alongside continuous measurements of HR (via three-lead ECG), pulse oximetry (SpO₂, Datex Ohmeda 3900, GE Healthcare, USA) and beat-to-beat measurements of SBP and DBP by photoplethysmography (Portapres, Finapres Medical Systems BV, The Netherlands). Change in SBP and DBP was calculated as the difference between pre-exercise and the value obtained at Watt_{max}. Urine specimens

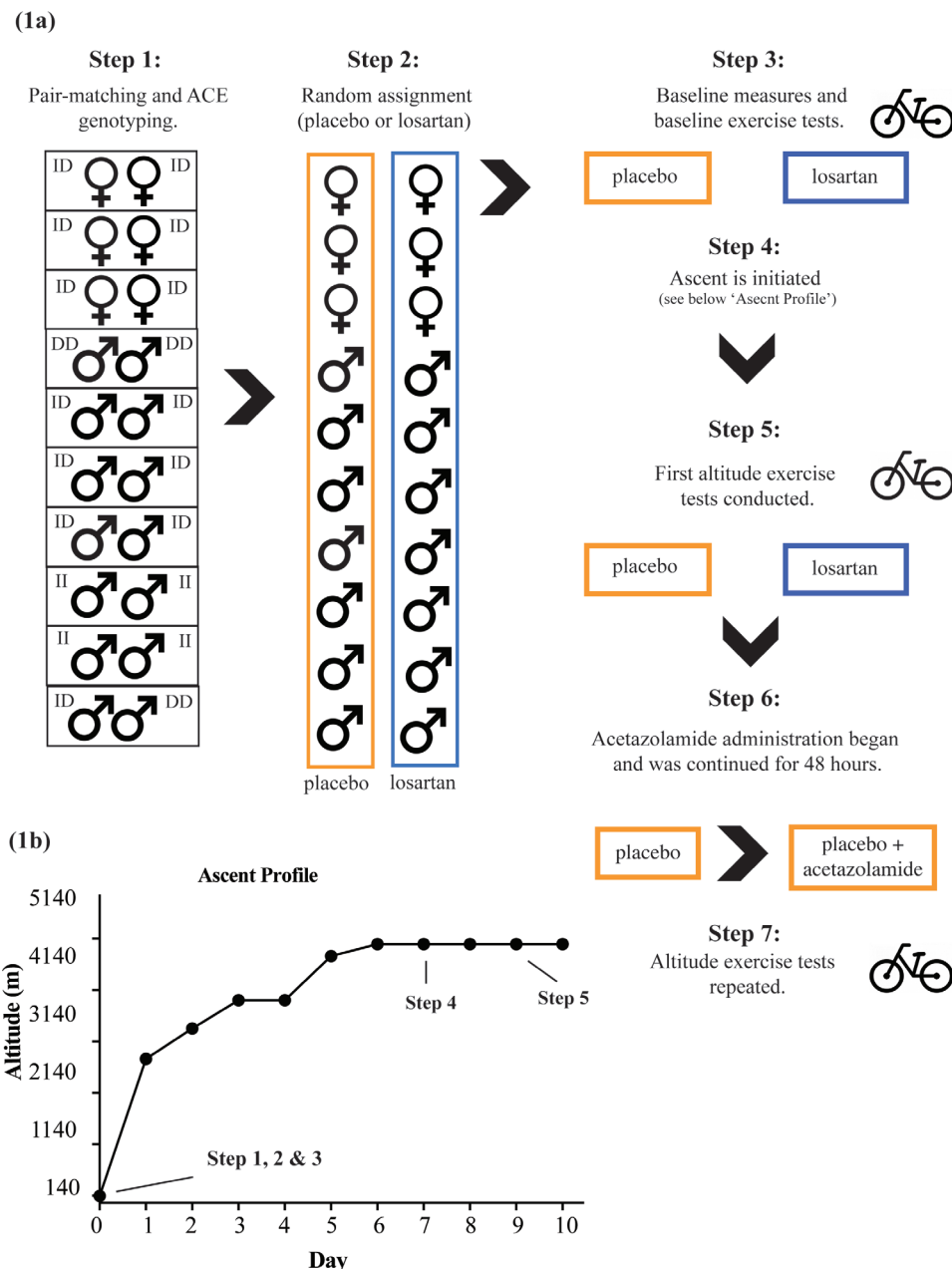


Figure 1 (A) Visual representation of the study design. Step 1: Participants were matched for: ACE genotype (II, ID or DD; see table 3), age, sex previous altitude exposure, and GFR. Step 2: Within each pair, participants were randomly assigned to placebo or losartan groups. Step 3: Baseline characteristics were recorded and baseline exercise tests were conducted (4 weeks before ascent) and were followed by the initiation of losartan administration (21 days before ascent). Step 4: Ascent is initiated with both groups ascending together in accordance with (B). Step 5: The first round of altitude exercise tests were conducted for members of both groups (5035 m). Step 6: Immediately following the first altitude exercise tests, acetazolamide was administered (125 mg orally, two times per day) and continued for 48 hours until next exercise test. Step 7: Repeat altitude exercise tests were conducted for all individuals (only placebo group data reported). (B) Expedition ascent profile. Day 0: Birmingham, UK (130 m), day 1: Quito, Ecuador (2800 m), day 2: Quito, Ecuador (2800 m), day 3: bus to Chunquiragua in Chaupi (3400 m), day 4: bus to Estrella del Chimborazo MARCO cruz (3950 m), day 5: Estrella del Chimborazo MARCO cruz (3950 m) with day hike to 5000 m and back, day 6: bus to Carrel hut (4800 m), days 7–10: Whympner hut (5035 m). GFR, glomerular filtration rate.

were collected surrounding exercise as outlined in the following section.

Urine collection and storage

Twenty-four-hour urine samples were collected over 1 day at sea level and on each day of the expedition (10 days)

with four aliquots (2 mL each) taken from each daily collection and frozen on dry ice.

For pre-exercise collections, participants were instructed to drink 500 mL of water between 90 and 30 min pre-exercise and to provide a timed (60 min) urine

specimen immediately prior to exercise. Post-exercise urine specimens were collected at 60, 120 and 180 min. Four aliquots (2 mL each) were taken from exercise specimens and frozen on dry ice, with the residual volumes returned to each individual's 24 hours collection bottle. Urine samples were transported back to Birmingham, UK and stored at (-80°C) until analysis.

Urine analysis

All samples were thawed at room temperature for 1 hour before analysis. The 24-hour urine samples were first analysed for $\alpha 1$ -AGP using radial immunodiffusion (RID; Talks et al., 2018)⁷ and the results were converted to excretion rates (mg/24 hours or $\mu\text{g}/\text{min}$). A subset of samples (119 out of 201) were then analysed using a latex-enhanced immunoassay on the Optilite turbidimetric analyser (The Binding Site, Birmingham, UK; online supplementary appendix 1). This automated method was used for all exercise samples, with the analysed concentrations (mg/L) converted to excretion rates ($\mu\text{g}/\text{min}$) based on sample volumes and collection durations.

728 29

Data analysis

Statistical analyses were performed using SPSS (IBM SPSS Statistics, V.25). Normality of distribution was determined by the Shapiro–Wilk test, with data (24-hour urinary $\alpha 1$ -AGP excretion) log-transformed where possible. Continuous variables are presented as mean \pm SD or median \pm IQR where appropriate. All tests for significance were two-tailed with statistical significance set at $p \leq 0.05$ unless otherwise indicated.

We used independent t-tests to compare group means (placebo vs losartan) where data were normally distributed and when measures were not repeated (eg, age and baseline HR). We used repeated-measures ANOVA with pairwise comparisons (Bonferroni corrected) to determine group (placebo vs losartan) and interaction effects (where appropriate) across days for normally distributed data (eg, transformed 24-hour $\alpha 1$ -AGP, DBP and SBP). We used Friedman tests to determine the main effect of time on $\alpha 1$ -AGP excretion surrounding exercise (ie, pre-60, post-60, post-120 and post-180 min) where data were not normally distributed. These results are presented as ' χ^2 (df), p value'. When appropriate, the post-hoc Wilcoxon signed rank test with Bonferroni correction for repeated measures was used to distinguish significance between time points (significance set at $p \leq 0.0125$; ie, corrected for three comparisons). We then used Mann–Whitney U tests to compare significant time points between groups (ie, placebo vs losartan at post-60 min).

To avoid any superimposed effects of the two drugs on results, we limited comparisons between the two altitude tests (before and after acetazolamide administration, placebo vs placebo +acetazolamide) to individuals from the placebo group completing both tests ($n=9$, tables 1 and 2). For these comparisons, we applied the Friedman

test and Wilcoxon signed rank test as previously described with additional use of the Wilcoxon signed rank test to compare $\alpha 1$ -AGP excretion rates (within-individuals) between placebo and placebo +acetazolamide at post-60 min. We used Spearman's correlation to evaluate relationships between daily measures (eg, DBP and 24 hours $\alpha 1$ -AGP excretion) and exercise measures (eg, SpO_2 and post-exercise $\alpha 1$ -AGP excretion).

We calculated ORs for urinary $\alpha 1$ -AGP excretion in placebo and losartan groups at rest (pre-exercise and post-exercise at 120 and 180 min) and with exercise (at post-60 min) based on the relative changes (relative increase (+) or decrease (–)) from baseline sea-level measures within each individual.

Patient and public involvement

This study was supported by the Birmingham Medical Research Expeditionary Society which provided input for the conduct of the research. Patients were not included. Public involvement was limited to recruitment. Notification was given to participants at the time of consent that acquisition of personal data was permitted on request. Permission was also obtained at this time for the dissemination of de-identified data within the research team and only externally when a reasonable request was submitted directly to the corresponding author(s) of the present study within 6 months of its publication. A portion of the cohort was invited to review the research methods for accuracy and readability.

Ethics approval

This study did not aim to investigate any safeties or efficacies of the already Food and Drug Administration (FDA)-approved drugs included, thus no clinical trial approval was obtained. There were no active FDA recalls for either drug for the duration of the study.

RESULTS

Baseline measures

Baseline data are presented in table 3. Placebo and losartan groups were not significantly different at baseline for measures of age, body mass, height, GFR, eGFR or creatinine. ACE genotype was exactly matched in 9 out of 10 pairs, with all allelic variations represented (ID, $n=13$; II, $n=4$ and DD, $n=3$; table 3).

Daily measures

Results for daily measures are presented in figure 2. Collectively with ascent, daily DBPs increased significantly ($p=0.04$) while SBP increased, although not significantly ($p=0.17$). Daily SBPs and DBPs remained similar between groups with ascent (SBP, $p=0.71$; DBP, $p=0.72$; figure 2A,B).

Exercise studies

Baseline sea-level exercise

Baseline exercise data are presented in table 1. All 20 participants achieved exhaustion during baseline sea-level exercise tests. Baseline measures of HR_{max} , absolute

Table 1 Maximal exercise test results compared between groups at baseline and twice at altitude

	Placebo	Losartan	P value		
Sea-level exercise					
HR _{max} (bpm)	171.7±16.5	173.7±22.9	0.82		
Absolute Watt _{max} (W)	243±66	238±70	0.89		
Relative Watt _{max} (W)	3.3±0.5	3.1±0.5	0.41		
	Placebo	Losartan	P value	Placebo +acetazolamide	P value
Altitude exercise					
<i>Pre-exercise</i>					
HR (bpm)	80±12	76±15	0.50	78±14	0.10
SBP (mm Hg)	162±28	159±22	0.82	149±22	0.17
DBP (mm Hg)	79±6	76±12	0.22	76±8	0.61
SpO ₂ (%)	76±7	78±5	0.40	84±4	<0.01*
<i>Maximal exercise</i>					
HR _{max} (bpm)	152±24	143±35	0.78	147±21	0.63
SBP at Watt _{max} (mm Hg)	229±47	221±50	0.81	232±47	0.43
ΔSBP (mm Hg)	70±31	66±42	0.80	82±46	0.26
DBP at Watt _{max} (mm Hg)	87±13	85±19	0.81	96±15	0.08
ΔDBP (mm Hg)	8±10	9±11	0.93	19±14	0.04 *
SpO ₂ at Watt _{max} (%)	70±5	71±7	0.89	77±4	<0.01*
Watt _{max} (W)	155±52	138±55	0.53	149±46	0.71
Watt _{max} (W/kg)	2.1±0.6	1.8±0.5	0.23	2.0±0.5	0.18

Values are presented as mean±SD. Change (Δ) SBP and ΔDBP represent the differences between pre-exercise and Watt_{max} measurements.

*Represents a significant (p≤0.05) difference between placebo and placebo +acetazolamide altitude exercise tests (n=9; within-individual comparisons).

DBP, diastolic blood pressure; HR, heart rate (bpm); HR_{max}, maximal heart rate; SBP, systolic blood pressure (mm Hg); SpO₂, oxygen saturation pulse oximetry (%); Watt_{max}, wattage at volitional fatigue (presented in absolute, W, and relative, W/kg, units).

Watt_{max} and relative Watt_{max} were not different between groups (table 1).

Altitude exercise (losartan vs placebo)

Altitude exercise data are presented in table 1. Pre-exercise measures of SpO₂, HR, SBP and DBP were not different between groups. All participants achieved exhaustion during the first altitude exercise tests. At altitude, both absolute Watt_{max} and relative Watt_{max} were reduced by ~40% compared with sea level but remained similar between groups. HR_{max}, SpO₂ at Watt_{max}, SBP at Watt_{max} and DBP at Watt_{max} were also similar between groups. Changes in SBP and changes in DBP did not differ between groups.

Altitude exercise (placebo vs placebo +acetazolamide)

Nine individuals from the placebo group went on to complete the second altitude exercise test following acetazolamide administration. These data are presented in table 1. Pre-exercise HR, SBP and DBP were no different between placebo and placebo + acetazolamide, although pre-exercise SpO₂ was significantly improved with acetazolamide (p<0.01).

Altitude exercise measures of absolute Watt_{max}, relative Watt_{max}, SBP at Watt_{max}, DBP at Watt_{max} and HR_{max} were similar before and after acetazolamide, as were the changes in SBP. In contrast, changes in DBP were significantly greater with acetazolamide (p=0.04). SpO₂ at Watt_{max} was significantly increased with acetazolamide (p<0.01).

Urine studies

Twenty-four-hour excretion

Twenty-four-hour urinary α1-AGP excretion (μg/min) significantly increased with ascent (p<0.01) although no differences were observed between groups for log-transformed excretion rates (p=0.97, figure 2). Twenty-four-hour urinary α1-AGP excretion correlated with DBP but not with SBP on days 1 (r=0.63, p=0.05), 6 (r=0.75, p=0.01) and 9 (r=0.90, p=0.04) of the expedition in the losartan group only.

Baseline sea level

Baseline urine results are presented in table 2 and figure 3A. Urinary α1-AGP excretion rates were no different between groups before exercise. Exercise increased urinary α1-AGP excretion in both placebo

Table 2 Comparisons of pre-exercise and post-exercise α 1-AGP excretion between groups at sea level and twice at altitude

	Placebo	P value (Z-score)	Losartan	P value (Z-score)		
Baseline α1-AGP excretion						
Pre-exercise	3.5 \pm 4.4	0.038 (–2.07)	1.4 \pm 2.8	0.028 (–2.20)		
Δ Pre-0 min to post-60 min	8.4 \pm 16.3	–	13.8 \pm 22.5	–		
Post-60 min	13.7 \pm 13.8	–	13.5 \pm 33.3	–		
Post-120 min	1.8 \pm 1.2	0.008 (–2.67)*	1.6 \pm 2.7	0.013 (–2.50)*		
Post-180 min	1.5 \pm 2.2	0.008 (–2.67)*	1.2 \pm 1.7	0.017 (–2.38)		
	Placebo	P value (Z-score)	Losartan	P value (Z-score)	Placebo +acetazolamide	P value (Z-score)
Altitude α1-AGP excretion						
Pre-exercise	3.1 \pm 5.3	0.022 (–2.29)	2.4 \pm 3.4	0.009 (–2.60)*	4.4 \pm 5.7	0.16 (–1.4)
Δ Pre- to post-60 min	7.9 \pm 14.3	–	4.8 \pm 9.9	–	1.2 \pm 11.03†	0.16 (–1.4)
Post-60 min	11.5 \pm 19.2	–	8.2 \pm 5.9	–	7.7 \pm 12.1	–
Post-120 min	3.2 \pm 2.4	0.007 (–2.70)*	2.8 \pm 4.4	0.012 (–2.52)*	3.4 \pm 5.0	–
Post-180 min	1.3 \pm 4.3	0.011 (–2.55)*	1.5 \pm 1.5	0.018 (–2.4)	2.7 \pm 4.2	0.025 (–2.24)

Urinary α 1-AGP excretion rates (μ g/min) are presented as median \pm IQR before (pre-exercise, 0 min) and after (post-60, post-120 and post-180 min) exercise initiation. Results are presented for placebo versus losartan (baseline), placebo versus losartan (first altitude exercise) and pre-acetazolamide versus acetazolamide (first compared with second altitude exercise).

*Represents the significance of post-60 min α 1-AGP excretion ($p \leq 0.0125$) compared with excretion at other time points.

†Represents the significant ($p \leq 0.05$) difference between groups (ie, placebo vs losartan or placebo vs placebo +acetazolamide) at the respective time point. Z-scores are presented in parenthesis where appropriate.

α 1-AGP, alpha-1 acid glycoprotein.

groups with excretion rates peaking at post-60 min (figure 3A). Urinary α 1-AGP excretion at post-60 min was similar between groups ($p=0.63$) and resolved after 120 min in both groups (figure 3A).

Altitude (placebo vs losartan)

Altitude exercise results are presented in table 2 and figure 3B. Collectively, pre-exercise α 1-AGP excretion was elevated at altitude compared with sea level, although excretion rates were similar between groups ($p=0.62$). The odds ratio (OR) for a relative increase in α 1-AGP

excretion at rest (pre-120, post-120, post-180 min) was 2.16 times greater without losartan at altitude.

Altitude exercise significantly increased urinary α 1-AGP excretion in both placebo ($\chi^2(3)=10.73$, $p=0.013$) and losartan ($\chi^2(3)=15.86$, $p<0.01$) groups (figure 3B). Post-60 min α 1-AGP excretion was lower with losartan compared with placebo, although the difference was not statistically significant ($p=0.28$, figure 3B and table 2).

In the losartan group only, the change in α 1-AGP excretion from pre-60 min to post-60 min was lower compared

Table 3 Baseline sea-level characteristics

	Placebo	Losartan	P value	ACE genotype and pair no.					
				Female pairs			Male pairs		
Age (years)	38.6 \pm 18.5	40.4 \pm 18.0	0.83	1	ID	ID	4	II	II
Body mass (kg)	74.1 \pm 11.5	66.7 \pm 13.3	0.71	2	ID	ID	5	II	II
Height (cm)	172.6 \pm 8.4	176.4 \pm 8.9	0.83	3	ID	ID	6	ID	ID
GFR (mL/min/1.73 m ²)	88.0 \pm 15.2	90.8 \pm 15.8	0.69				7	ID	ID
eGFR (mL/min/1.73 m ²)	87.8 \pm 15.3	90.8 \pm 15.9	0.68				8	ID	ID
Creatinine (μ mol/L)*	84.5 \pm 24.0	81.0 \pm 31.0	0.58				9	DD	ID
							10	DD	DD

Baseline results were obtained at sea level and prior to the administration of placebo and losartan. Results for continuous variables are reported as mean \pm SD unless indicated by *, where values are reported as median \pm IQR. ACE genotypes are presented as allelic variations (II, ID and DD). Significance was set to p value <0.05 . No significant differences between groups were evident at baseline sea level. Subjects were equally matched for sex (men and women). All genotypes were observed and 90% matched.

eGFR, estimated glomerular filtration rate (calculated using the MDRD study equation); GFR, glomerular filtration rate; MDRD, Modification of Diet in Renal Disease.

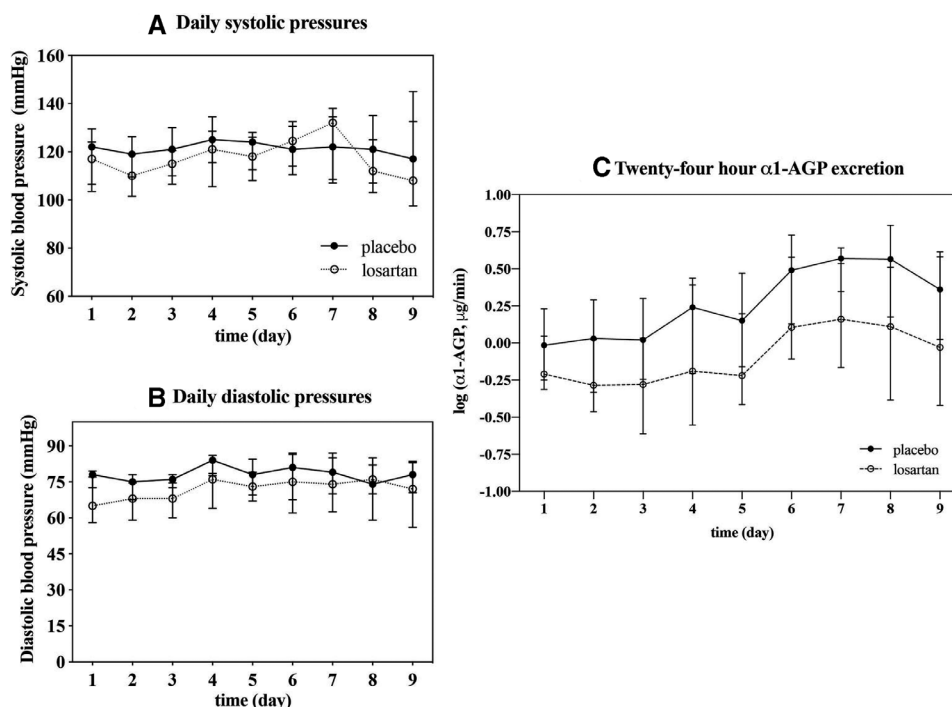


Figure 2 Daily measures of blood pressure and 24-hour urinary α 1-AGP excretion rates with ascent. (A) Daily SBPs (mm Hg). (B) Daily DBPs (mm Hg). Data are plotted as daily medians with error bars representing the respective IQRs. (C) Log-transformed 24-hour urinary α 1-AGP excretion rates (μ g/min) by days with ascent. Data are plotted as the mean log of 24 hours α 1-AGP with error bars representing SD of the respective group mean on each day. Twenty-four-hour urinary α 1-AGP excretion collectively increased with ascent ($p<0.01$) with no difference between groups ($p=0.97$). α 1-AGP, alpha-1 acid glycoprotein; DBPs, diastolic blood pressures; SBPs, systolic blood pressures.

with those changes observed at baseline, although this difference was not significant ($p=0.059$, figure 3D). The OR for reduced urinary α 1-AGP excretion at post-60 min (sea level vs first altitude test) was 2.25 times greater with losartan.

Altitude (placebo vs placebo +acetazolamide)

Results for comparisons of post-exercise α 1-AGP between placebo and placebo +acetazolamide are presented in tables 1 and 2 and figure 3C,E. As with placebo, altitude exercise tests, placebo +acetazolamide exercise resulted in an increase in urinary α 1-AGP excretion ($\chi^2(3)=10.73$, $p=0.01$, figure 3C). Despite similar exercise peak power outputs (table 1) between placebo and placebo +acetazolamide tests, post-exercise α 1-AGP excretion (at 60 min post-exercise) was significantly lower following placebo +acetazolamide exercise ($p=0.025$, figure 3C). Exercise-induced increases in α 1-AGP excretion (change from pre-exercise to post-60 min) were significantly reduced ($p=0.036$) with acetazolamide by nearly threefold (figure 3E). These changes were not correlated ($r=-0.10$, $p=0.82$) with the significant improvements in pre-exercise SpO_2 or SpO_2 at $Watt_{max}$ with acetazolamide administration (table 2). No difference was observed between losartan and losartan +acetazolamide tests (data not shown).

DISCUSSION

Compared with sea level and despite substantial systemic hypoxia at $Watt_{max}$, post-exercise α 1-AGP excretion was

not greater at altitude suggesting that hypoxia is not the primary mechanism. Altitude-related reductions in exercise intensity could, in part, explain this result,³⁰ but they would fail to explain the increased likelihood for post-exercise α 1-AGP to be lower with losartan compared with placebo when exercise intensities were similar. We had expected that losartan would mitigate BP amplifications and thus lower post-exercise α 1-AGP, however we observed no difference in the BP response to exercise between groups. Therefore, we have no evidence to attribute post-exercise α 1-AGP responses to alterations in³¹BP or peritubular pressures.³²

The direct action of ARBs within the glomerular filtration barrier on ATIs of podocytes¹⁶ could provide an alternative explanation. Activation of ATIs on podocytes induces heparanase expression, which promotes the cleavage of heparan sulfate and inhibits the production and secretion of heparan sulfate proteoglycans (HSPGs).³³ The net result is neutralisation of the negatively charged HSPGs which limits the charge-selective function of the glomerular basement membrane (GBM).³⁴ Blocking ATIs with losartan thus promotes the localised retention of negatively charged proteins at the GBM³⁵ and limits the charge-selectivity impairment³⁶ which, in the present study, manifests as a reduction in the extent of post-exercise α 1-AGP. This hypothesis is consistent with findings related to the intensity-dependent increases in angiotensin-II³⁷ and

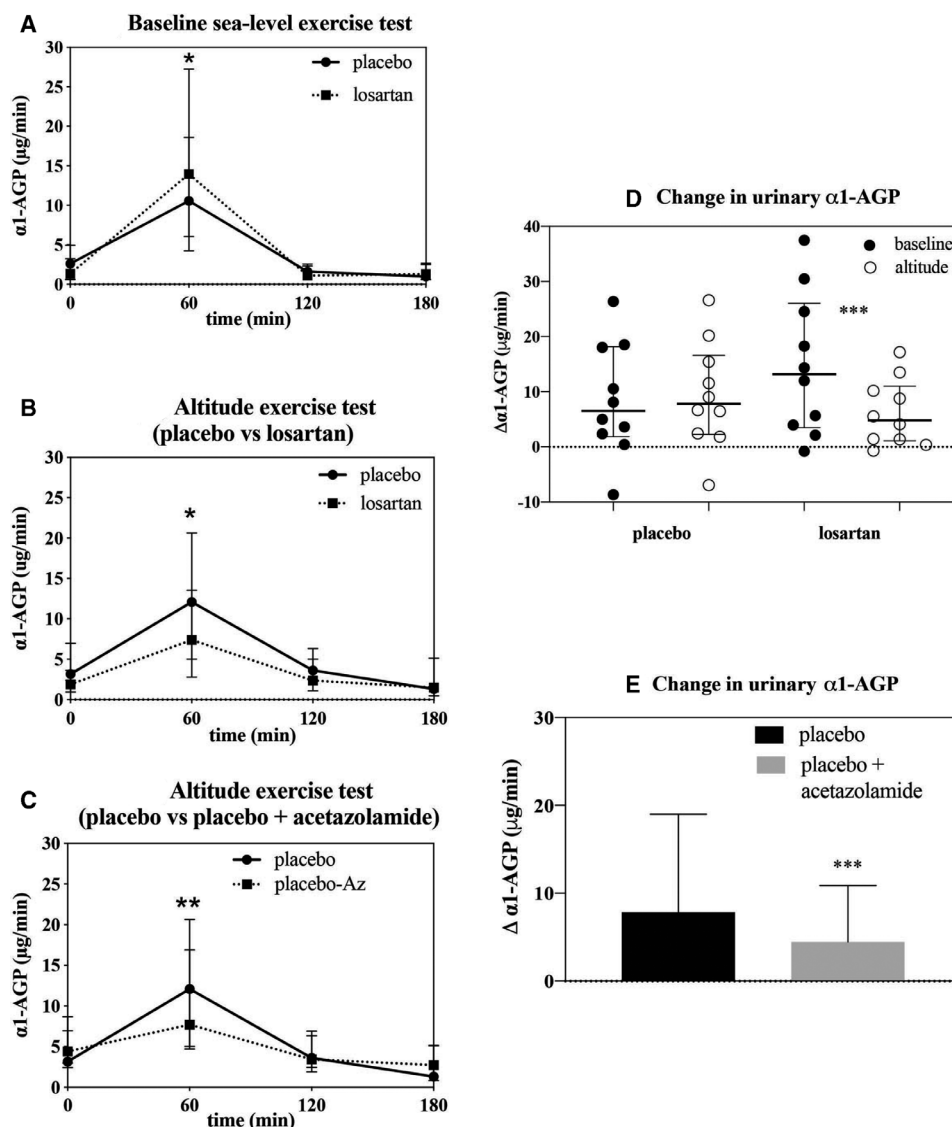


Figure 3 Pre-exercise and post-exercise urinary $\alpha 1$ -AGP excretion ($\mu\text{g}/\text{min}$) rates. (A) Comparisons between placebo versus losartan groups at baseline sea level; (B) comparisons between placebo versus losartan groups at altitude (first altitude exercise); (C) comparisons between placebo versus placebo + acetazolamide at altitude (second altitude exercise); (D) comparisons between baseline versus altitude in both placebo and losartan groups (change in $\alpha 1$ -AGP from pre-60 min to post-60 min, $\Delta\alpha 1$ -AGP) and (E) comparisons between placebo versus placebo + acetazolamide. Results are plotted as the group median (or individual values, D) with error bars representing the relative IQRs of the group. Significance was set to p value ≤ 0.05 unless otherwise indicated. Representing significance: * for the significant effect of exercise on urinary $\alpha 1$ -AGP excretion; ** for the significant difference between groups at post-60 min and *** for the significant difference between placebo and placebo + acetazolamide for $\Delta\alpha 1$ -AGP (D) or for the trend ($p=0.059$) of difference between baseline and the first altitude for $\Delta\alpha 1$ -AGP (E). $\alpha 1$ -AGP, alpha-1 acid glycoprotein.

exercise-related increases in $\alpha 1$ -AGP,³⁸ although further investigations are required.

Reductions in post-exercise $\alpha 1$ -AGP by acetazolamide were unrelated to the significant improvements in exercise SpO_2 at similar intensity (these effects of acetazolamide on exercise performance were previously published and support previous findings).^{26–39} On its own, this would provide further support indicating that hypoxia is not the primary mechanism of post-exercise proteinuria. Unfortunately, the post-exercise effects of acetazolamide cannot be separated from the acclimatisation effect. Thus, no definitive conclusion regarding

acetazolamide's effects on post-exercise proteinuria can be provided. Future research is required to evaluate acetazolamide-related changes in post-exercise proteinuria. Nonetheless, acetazolamide results support recent evidence demonstrating performance limiting effects of acetazolamide despite of its ability to elicit improvements in oxygen saturation.⁴⁰

Limitations and future directions

The inability to control for extraneous variables (eg, activity, sleep, environmental conditions) as well as low subject numbers were weaknesses of the present study

but they are common limitations for such field-based research. Not controlled for were the inter-individual variability⁷ and between-day variations in $\alpha 1$ -AGP excretion which could have impacted results. Future research could be strengthened by incorporating a sea-level control arm that compared post-exercise proteinuria between exercise tests executed 48 hours apart. Benefit could also be gained by using a less complex design and avoiding overlapping factors (eg, acclimatisation status and time course of exercise tests in this case). Lastly, further research is needed in order to confirm the relationship between DBP and 24 hours $\alpha 1$ -AGP excretion.

CONCLUSION

Our findings suggest that post-exercise $\alpha 1$ -AGP is (1) more related to exercise intensity than degree of hypoxia or BP and (2) possibly influenced by activation or inhibition of AT1 receptors. Losartan may limit post-exercise proteinuria by helping to maintain the charge-selectivity function in the glomerular filter, although further investigations are required for evaluation.

Author affiliations

¹School of Sport, Exercise, and Rehabilitation Sciences, University of Birmingham, Birmingham, UK

²Birmingham Medical Research Expeditionary Society, University of Birmingham, Birmingham, UK

³Institute of Clinical Sciences, University of Birmingham, Birmingham, UK

⁴Medical School, East Surrey Hospital, Redhill, UK

⁵Occupational Performance Research Group, University of Chichester Department of Sport and Exercise Sciences, Chichester, UK

⁶Research & Development, Binding Site Group Ltd, Edgbaston, UK

⁷St. George's University Hospital, University of London, London, UK

⁸School of Biological Sciences, University of Portsmouth, Portsmouth, UK

⁹Department of Anaesthesia, Ninewells Hospital, Dundee, UK

¹⁰Vascular Surgery, University Hospitals Coventry and Warwickshire NHS Trust, Coventry, UK

¹¹Department of Bioengineering, Imperial College London, London, UK

¹²Academic Foundation Programme, NHS Highland, Inverness, United Kingdom

¹³University Hospitals Birmingham NHS Foundation Trust, Birmingham, UK

¹⁴BASEM, Doncaster, UK

¹⁵Medical School, University of Birmingham, Birmingham, UK

Acknowledgements The authors would like to thank Gregg Wallace and the Research and Development Team at The Binding Site for their kind assistance during the development of the immunoassay and subsequent urinalysis. The authors would also like to thank the Birmingham Medical Research Expeditionary Society.

Contributors All authors listed contributed substantially to the work and by ICMJE 2018 standards meet at minimum satisfy criterion No. 1, No. 3 and No. 4 with additional support for No. 2 provided by KEJ, SJEL, AF, AW, JD and AB.

Funding The research was partially funded by JABBS Foundation.

Competing interests None declared.

Patient and public involvement Patients and/or the public were not involved in the design, conduct, reporting or dissemination plans of this research. Refer to the Methods section for further details.

Patient consent for publication Not required.

Ethics approval This observational study was approved by the Chichester University Research Ethics Committee (protocol number: 1314 42) and was performed according to the Declaration of Helsinki.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request. The data are owned by the University of Birmingham and can be obtained upon reasonable

request from either the corresponding author or the Sport, Exercise, and Rehabilitation Sciences department +44 (0)121 414 9286. Reuse is not permitted unless otherwise indicated at the time of reasonable request. There is no additional relevant information.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iDs

Kelsley E Joyce <http://orcid.org/0000-0002-3189-847X>

Arthur Bradwell <http://orcid.org/0000-0002-1562-1606>

REFERENCES

- Winterborn MH, Bradwell AR, Chesner IM, *et al.* The origin of proteinuria at high altitude. *Postgrad Med J* 1987;63:179–81.
- Poortmans JR. Exercise and renal function. *Sports Med* 1984;1:125–53.
- Christensen EI, Birn H, Storm T, *et al.* Endocytic receptors in the renal proximal tubule. *Physiology* 2012;27:223–36.
- Jarad G, Miner JH. Update on the glomerular filtration barrier. *Curr Opin Nephrol Hypertens* 2009;18:226–32.
- Poortmans JR. Postexercise proteinuria in humans. Facts and mechanisms. *JAMA* 1985;253:236–40.
- Poortmans J, Jeanloz RW. Quantitative immunological determination of 12 plasma proteins excreted in human urine collected before and after exercise. *J Clin Invest* 1968;47:386–93.
- Talks BJ, Bradwell SB, Delamere J, *et al.* Urinary alpha-1-acid glycoprotein is a sensitive marker of glomerular protein leakage at altitude. *High Alt Med Biol* 2018;19:295–8.
- Jerebtsova M, Saraf SL, Soni S, *et al.* Urinary orosomucoid is associated with progressive chronic kidney disease stage in patients with sickle cell anemia. *Am J Hematol* 2018;93:E107–9.
- Christiansen MS, Hommel E, Magid E, *et al.* Orosomucoid in urine is a powerful predictor of cardiovascular mortality in normoalbuminuric patients with type 2 diabetes at five years of follow-up. *Diabetologia* 2005;48:386–93.
- Wolyniec W, Kasprowicz K, Rita-Tkachenko P, *et al.* Biochemical markers of renal hypoperfusion, hemoconcentration, and proteinuria after extreme physical exercise. *Medicina* 2019;55. doi:10.3390/medicina55050154. [Epub ahead of print: 17 May 2019].
- Bradwell A, Brearey S, Harris S, *et al.* Effect of hard exercise on proteinuria at high altitude. *High Alt Med Biol* 2003;3:436.
- Luks AM, Johnson RJ, Swenson ER. Chronic kidney disease at high altitude. *J Am Soc Nephrol* 2008;19:2262–71.
- Caravita S, Faini A, Baratto C, *et al.* Upward shift and steepening of the blood pressure response to exercise in hypertensive subjects at high altitude. *J Am Heart Assoc* 2018;7:1–7.
- Bilo G, Villafuerte FC, Faini A, *et al.* Ambulatory blood pressure in untreated and treated hypertensive patients at high altitude: the high altitude cardiovascular Research-Andes study. *Hypertension* 2015;65:1266–71.
- Parati G, Bilo G, Faini A, *et al.* Changes in 24 h ambulatory blood pressure and effects of angiotensin II receptor blockade during acute and prolonged high-altitude exposure: a randomized clinical trial. *Eur Heart J* 2014;35:3113–22.
- Xu F, Mao C, Liu Y, *et al.* Losartan chemistry and its effects via AT1 mechanisms in the kidney. *Curr Med Chem* 2009;16:3701–15.
- Koeppen BM, Stanton BA. *Renal physiology*. 5th edn. Philadelphia, PA: Elsevier Mosby, 2013.
- West JB, Lahiri S, Gill MB, *et al.* Arterial oxygen saturation during exercise at high altitude. *J Appl Physiol* 1962;17:617–21.
- Jonk AM, van den Berg IP, Olfert IM, *et al.* Effect of acetazolamide on pulmonary and muscle gas exchange during normoxic and hypoxic exercise. *J Physiol* 2007;579:909–21.
- Joyce KE, Lucas SJE, Imray CHE, *et al.* Advances in the available non-biological pharmacotherapy prevention and treatment of acute mountain sickness and high altitude cerebral and pulmonary oedema. *Expert Opin Pharmacother* 2018;19:1891–902.
- Leaf DE, Goldfarb DS. Mechanisms of action of acetazolamide in the prophylaxis and treatment of acute mountain sickness. *J Appl Physiol* 2007;102:1313–22.
- Bradwell AR, Delamere JP. The effect of acetazolamide on the proteinuria of altitude. *Aviat Space Environ Med* 1982;53:40–3.
- Parving H-H, de Zeeuw D, Cooper ME, *et al.* ACE gene polymorphism and losartan treatment in type 2 diabetic patients with nephropathy. *J Am Soc Nephrol* 2008;19:771–9.

- 24 Bigham AW, Kiyamu M, León-Velarde F, *et al.* Angiotensin-converting enzyme genotype and arterial oxygen saturation at high altitude in Peruvian Quechua. *High Alt Med Biol* 2008;9:167–78.
- 25 Levey AS, Bosch JP, Lewis JB, *et al.* A more accurate method to estimate glomerular filtration rate from serum creatinine: a new prediction equation. modification of diet in renal disease Study Group. *Ann Intern Med* 1999;130:461–70.
- 26 Bradwell AR, Myers SD, Beazley M, *et al.* Exercise limitation of acetazolamide at altitude (3459 M). *Wilderness Environ Med* 2014;25:272–7.
- 27 Thompson PD, Arena R, Riebe D, *et al.* ACSM's new preparticipation health screening recommendations from ACSM's guidelines for exercise testing and prescription, ninth edition. *Curr Sports Med Rep* 2013;12:215–7.
- 28 Kustán P, Szirmay B, Horváth-Szalai Z, *et al.* Urinary orosomucoid: validation of an automated immune turbidimetric test and its possible clinical use. *Biochem Med* 2016;26:421–30.
- 29 Christiansen MS, Blirup-Jensen S, Foged L, *et al.* A particle-enhanced turbidimetric immunoassay for quantitative determination of orosomucoid in urine: development, validation and reference values. *Clin Chem Lab Med* 2004;42:1168–77.
- 30 Poortmans JR, Labilloy D. The influence of work intensity on postexercise proteinuria. *Eur J Appl Physiol Occup Physiol* 1988;57:260–3.
- 31 Rangemark C, Lind H, Lindholm L, *et al.* Lisinopril reduces postexercise albuminuria more effectively than atenolol in primary hypertension. *Eur J Clin Pharmacol* 1996;49:267–71.
- 32 Esnault VL, Potiron-Josse M, Testa A, *et al.* Captopril but not acebutolol, prazosin or indomethacin decreases postexercise proteinuria. *Nephron* 1991;58:437–42.
- 33 Kramer A, van den Hoven M, Rops A, *et al.* Induction of glomerular heparanase expression in rats with adriamycin nephropathy is regulated by reactive oxygen species and the renin–angiotensin system. *J Am Soc Nephrol* 2006;17:2513–20.
- 34 Brinkkoetter P-T, Holtgrete S, van der Woude FJ, *et al.* Angiotensin II type 1-receptor mediated changes in heparan sulfate proteoglycans in human SV40 transformed podocytes. *J Am Soc Nephrol* 2004;15:33–40.
- 35 Deynelli O, Yavuz D, Velioglu A, *et al.* Effects of ACE inhibition and angiotensin II receptor blockade on glomerular basement membrane protein excretion and charge selectivity in type 2 diabetic patients. *J Renin Angiotensin Aldosterone Syst* 2006;7:98–103.
- 36 van Det NF, Tamsma JT, van den Born J, *et al.* Differential effects of angiotensin II and transforming growth factor beta on the production of heparan sulfate proteoglycan by mesangial cells in vitro. *J Am Soc Nephrol* 1996;7:1015–23.
- 37 van Ginkel S, de Haan A, Woerdeman J, *et al.* Exercise intensity modulates capillary perfusion in correspondence with ACE I/D modulated serum angiotensin II levels. *Appl Transl Genom* 2015;4:33–7.
- 38 Joyce KE, Balanos GM, Fountain A, *et al.* Hypoxia does not influence post-exercise proteinuria. *European Database of Sport Science* 2019. (abstract).
- 39 Bradwell AR, Ashdown K, Rue C, *et al.* Acetazolamide reduces exercise capacity following a 5-day ascent to 4559 M in a randomised study. *BMJ Open Sport Exerc Med* 2018;4:e000302.
- 40 Elisabeth E, Hannes G, Johannes B, *et al.* Effects of low-dose acetazolamide on exercise performance in simulated altitude. *Int J Physiol Pathophysiol Pharmacol* 2017;9:28–34.